

PRE-APPEAL BRIEF REQUEST FOR REVIEW

Docket Number (Optional)

2916693-014000

I hereby certify that this correspondence is being deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to "Mail Stop AF, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450" [37 CFR 1.8(a)]

on _____

Signature _____

Typed or printed name _____

Application Number

10782096

Filed

February 19, 2004

First Named Inventor

Carozzi et al.

Art Unit

1638

Examiner

KUBELIK, Anne R.

Applicant requests review of the final rejection in the above-identified application. No amendments are being filed with this request.

This request is being filed with a notice of appeal.

The review is requested for the reason(s) stated on the attached sheet(s).

Note: No more than five (5) pages may be provided.

I am the

☐ applicant/inventor.

/David L. Vanik/

Signature

☐ assignee of record of the entire interest.
See 37 CFR 3.71. Statement under 37 CFR 3.73(b) is enclosed.
(Form PTO/SB/96)

David L. Vanik

Typed or printed name

☒ attorney or agent of record.
Registration number 64,547

202-508-3400

Telephone number

☐ attorney or agent acting under 37 CFR 1.34.

July 12, 2010

Date

Registration number if acting under 37 CFR 1.34 _____

NOTE: Signatures of all the inventors or assignees of record of the entire interest or their representative(s) are required.
Submit multiple forms if more than one signature is required, see below.

☒ *Total of 3 forms are submitted.

This collection of information is required by 35 U.S.C. 132. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11, 1.14 and 41.6. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Mail Stop AF, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

If you need assistance in completing the form, call 1-800-PTO-9199 and select option 2.

Privacy Act Statement

The **Privacy Act of 1974 (P.L. 93-579)** requires that you be given certain information in connection with your submission of the attached form related to a patent application or patent. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) furnishing of the information solicited is voluntary; and (3) the principal purpose for which the information is used by the U.S. Patent and Trademark Office is to process and/or examine your submission related to a patent application or patent. If you do not furnish the requested information, the U.S. Patent and Trademark Office may not be able to process and/or examine your submission, which may result in termination of proceedings or abandonment of the application or expiration of the patent.

The information provided by you in this form will be subject to the following routine uses:

1. The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether disclosure of these records is required by the Freedom of Information Act.
2. A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
3. A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
4. A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552a(m).
5. A record related to an International Application filed under the Patent Cooperation Treaty in this system of records may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant to the Patent Cooperation Treaty.
6. A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or his/her designee, during an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 122(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspection or an issued patent.
9. A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re the application of:

Confirmation No.: 5854

Carozzi *et al.*

Group Art Unit: 1638

Application Serial No.: 10/782,096

Examiner: Anne R. Kubelik

Filed: February 19, 2004

Attorney Docket No.: 2916693-014000

For: AXMI-009, A DELTA-ENDOTOXIN GENE AND METHODS FOR ITS USE

Pre-Appeal Brief Request for Review

Dear Members of the Panel:

The enclosed Pre-Appeal Brief Request is filed along with a Notice of Appeal dated July 12, 2010 and is accompanied by payment of a three-month extension of time request and fee.

Pending Claims

Claims 1-11, 19, and 22-23 are pending in this application of which Claims 1, 22, and 23 are independent.

Pending Rejections

Claims 1 and 4-7 stand rejected under 35 U.S.C. § 103(a) as being allegedly unpatentable over Ben-Dov *et al.* (*Appl. Environ. Microbiol.* 62, pages 3140-3145, 1996) in view of Carlton *et al.* (*Mol. Biol. Microb. Differ., Proc. Intl. Spore Conf.*, 9th, Meeting date 1984, pages 246-252, 1985) and further in view of Applicant's response to the Request for Information under 37 C.F.R. 1.105. Claims 2-3, 8-11, 19, 22-23 stand rejected under 35 U.S.C. § 103(a) as being allegedly unpatentable over Ben-Dov *et al.* in view of Carlton *et al.* and Koziel *et al.* (U.S. Patent No. 5,625,136). Applicants respectfully disagree and request withdrawal of the rejections.

- (1) **Rejection of Claims 1 and 4-7 under 35 U.S.C. § 103(a) over Ben-Dov *et al.* in view of Carlton *et al.* and in view of Applicant's response to the Request for Information under 37 C.F.R. 1.105**

- (a) One of ordinary skill in the art would have no motivation to combine Ben-Dov *et al.* together with Carlton *et al.*

At the outset, the Examiner impermissibly uses Applicant's Specification as a basis for the rejection. Specifically, the Examiner requested source information under 37 C.F.R. 1.105 and

subsequently used this information as a basis for the obviousness rejection under 35 U.S.C. § 103(a).¹ However, outside of Applicant's Specification, one of ordinary skill in the art would have no reason to select and isolate sequences from HD536 given the numerous possibilities of well-known strains exhibiting insecticidal activity. Moreover, the only motivation for even considering sequences isolated from HD536 comes from the instant disclosure and not the cited references. Without a specific teaching or motivation, one of ordinary skill in the art would not even look to Carlton *et al.* to choose a strain, let alone select HD536 from the laundry list of possibilities included in Carlton *et al.*

Ben-Dov *et al.* does not teach or suggest all of the claimed elements. The Examiner acknowledges that Ben-Dov *et al.* alone is deficient to render the claims obvious and states that "Ben-Dov *et al.* do not teach a nucleic acid encoding SEQ ID NO: 2, 4 or 6." Final Office Action at page 4. At best, Ben-Dov *et al.* teach cloning of large restriction fragments from *Bacillus thuringiensis* subsp. *israelensis* and identification of known toxins using Southern hybridization and probes specific for the known toxins. Ben-Dov *et al.* does not teach or suggest a nucleic acid molecule encoding polypeptide with activity against lygus pests, let alone the claimed sequences. Rather, Ben-Dov *et al.* is concerned with characterizing a single 125-kilobase plasmid containing genes that encode delta-endotoxins with activity against mosquito larvae. Ben-Dov *et al.* at page 3140.

Carlton *et al.* evaluates a large number of *Bacillus thuringiensis* strains for the presence of extrachromosomal DNA by agarose gel electrophoresis. Carlton *et al.* at page 251 and Figure 1. In doing so, Carlton *et al.* sets forth a laundry list of *Bacillus thuringiensis* plasmids, including HD536, together with the estimated size of the plasmids in Table 1. Carlton *et al.* at Table 1. In rejecting the claims, the Examiner cites to Table 1 and states that "Carlton *et al.* teach that strain HD536 has a 68 MDa plasmid implicated in toxin production." *Id.* The Examiner further asserts that "it would have been obvious to one of ordinary skill in the art to modify the method of cloning delta-endotoxin genes from *B. thuringiensis* plasmids as taught by Ben-Dov *et al.*, to clone delta-endotoxin genes from strain HD536 described in Carlton *et al.*" *Id.* The Examiner also states that "[o]ne of ordinary skill in the art would have been motivated to do so because an increased

¹ Applicants respectfully note that the response to the Request for Information under 37 CFR 1.105 was submitted to the USPTO and was labeled as proprietary material with the label "DO NOT SCAN." Applicants respectfully disagree with the Examiner's inclusion of this material in the Final Office Action dated January 12, 2010.

repertoire of delta-endotoxins would be desirable for increasing toxicity spectra and for overcoming pest resistance to existing endotoxins.” *Id.* Applicants respectfully disagree.

One of ordinary skill in the art would have no motivation to combine the teachings of Ben Dov *et al.* with Carlton *et al.* in a manner that renders the claims obvious. For one, Ben Dov *et al.* is limited to evaluating a specific 125-kilobase plasmid containing known genes that encode delta-endotoxins with activity against mosquito larvae. Ben-Dov *et al.* at page 3140. To achieve this, Ben-Dov *et al.* utilizes probes capable of specifically identifying genes of interest that encode delta-endotoxins with specific activity against mosquito larvae. Ben-Dov *et al.* provides no additional motivation to apply this methodology to toxins other than those that are active against mosquito larvae. Ben-Dov *et al.* also acknowledges that “[a]mbiguous results were obtained with several additional probes (data not shown)...”, thus indicating further uncertainty regarding the methodology and applicability of the teachings of Ben-Dov *et al.* Given the above, one of ordinary skill in the art would have no motivation to modify and apply the methodology of Ben-Dov *et al.* to other plasmids, such as those set forth in Carlton *et al.*, with a reasonable expectation of success.

An “obvious to try” rationale may only support a conclusion that a claim would have been obvious where one skilled in the art is choosing from a finite number of identified, predictable solutions, with a reasonable expectation of success. *KSR Int’l Co. v. Tele-flex Inc.*, 550 U.S. 389 (2007). One of ordinary skill in the art would recognize that there was no reasonable expectation of success in obtaining any toxin genes from HD536 since no insecticidal activity was demonstrated for this strain prior to the Applicant’s disclosure. Outside of providing a loose correlation between the ~68 MDa plasmid of HD536 and “toxin production,” Carlton *et al.* fails to suggest that genes isolated from HD536 would have any insecticidal activity. Instead, Carlton *et al.* merely suggest that the 68 MDa plasmid present in strain HD536 may be responsible for crystal protein production. The presence of a crystal protein provides no evidence for the presence of a gene or encoded protein having insecticidal activity against any pest, particularly any lepidopteran, coleopteran or heteropteran pests.

(b) One of ordinary skill in the art would have no motivation to use the specific probes of Ben-Dov *et al.* to isolate SEQ ID NO:1-6 from HD536

In rejecting the claims, the Examiner cites to Carlton *et al.* and states that “knowledge that the 68 KDa plasmid encodes a toxins would motivate one of skill in the art to sequence the plasmid

to search for the toxins genes.” Final Office Action at page 5. Applicants respectfully disagree. One of ordinary skill in the art would have no motivation to use the methodology and specific probes of Ben-Dov *et al.* in an attempt to isolate any one of claimed SEQ ID NO:1-6 from HD536. For one, the claimed sequences have a low sequence homology with other known toxins (<30%). As such, one of skill would not have used the cryIVA, cryIVB, cryIVC, cryIVD, and cytA, probes taught in Ben-Dov *et al.* to isolate any sequence from HD536, let alone the claimed sequences. In fact, only the cryIVA probe was able to detect any gene other than itself, and Ben-Dov *et al.* appears to attribute this cross-reactivity with the degree of sequence homology between the two genes. See Ben-Dov *et al.* at column 2, page 3143. Outside of pure conjecture, it is not clear how one of skill in the art would be able to use the hybridization method disclosed by Ben-Dov *et al.* to isolate the sequences of the invention.

(c) One of ordinary skill in the art would have no motivation to try and isolate the claimed sequences from HD536 of Carlton *et al.*

A person of ordinary skill in the art would have no motivation to specifically select and isolate sequences from HD536, especially given the laundry list of plasmids described in Carlton *et al.* Additionally, HD536 is only mentioned once in Table 1 and Carlton *et al.* does not specifically indicate why it would be advantageous to isolate sequences from HD536, let alone the claimed sequences. As there is no structural similarity between the known cry toxins identified by Ben-Dov *et al.* and the claimed sequences, a person of ordinary skill in the art would not looked to the methodology of Ben-Dov *et al.* in an attempt to isolate the sequences of the instant invention from HD536.

(d) AXMI-007 unexpectedly exhibits insecticidal activity against *lygus lineolaris*

For at least the reasons set forth above, Applicants submit that the Examiner has failed to provide a *prima facie* case of obviousness. However, irrespective of this, secondary considerations of the advantageous properties of the claimed sequences, particularly the broad insecticidal activities of the recited sequences, provide additional support for the nonobviousness of the pending claims. For instance, Example 11 of the Specification provides evidence of the insecticidal efficacy of AXMI-009 against *lygus lineolaris*. Specification, for example, at page 39, line 6 – page 40, line 2. As set forth in Example 11, samples containing the AXMI-009 protein in *Bacillus* yield a mortality rate of 50% against *lygus lineolaris* relative to a 0% mortality rate for the control.

Specification at Table 4. This result is particularly unexpected given the relatively low amino acid identity of AXMI-009 as compared to the 17 exemplary endotoxin classes described in Table 1 of the Specification (<30%).

(2) Rejection of Claims 2-3, 8-11, 19, 22-23 under 35 U.S.C. § 103(a) over Ben-Dov *et al.* in view of Carlton *et al.* and Koziel *et al.*

In rejecting Claims 2-3, 8-11, 19, 22-23 under 35 U.S.C. § 103(a), the Examiner acknowledges that “Ben-Dov *et al.* in view of Carlton *et al.* do not teach plants and seeds transformed with the nucleic acid.” Final Office Action at page 7. However, the Examiner cites to Koziel *et al.* and asserts that “[a]t the time the invention was made, it would have been obvious to one of ordinary skill in the art to transform the nucleic acid taught by Ben-Dov *et al.* in view of Carlton *et al.* into plants, including maize, as described in Koziel *et al.*” *Id.*

For at least the reasons set forth above, Applicants respectfully disagree with the Examiner's rejection of Claims 3, 8-11, 19, 22-23. Moreover, one of ordinary skill in the art would have no motivation to transform the nucleic acids taught by Ben-Dov *et al.* in view of Carlton *et al.* into plants or cells, let alone the specific claimed plants and cells. For one, Ben-Dov *et al.* has no association with plants and is solely concerned with isolating genes that encode toxins that are active against mosquito larvae. Ben-Dov *et al.* at page 3140. Accordingly, the emphasis in Ben-Dov *et al.* is centered around human infectious diseases as compared to plants or plant cells. *Id.* Outside of including HD536 in a list and describing a loose correlation between HD536 and toxin activity, Carlton *et al.* provides no motivation for including SEQ ID NO: 1, 2, 3, or 4 in plants or cells. Koziel *et al.* fails to remedy the deficiencies of both Ben-Dov *et al.* and Carlton *et al.* and does not suggest transforming the claimed sequences into plants or cells.

For at least the above, withdrawal of the rejections is respectfully requested.

July 12, 2010

Customer No.: **95725**

Telephone: 202-508-3400

Respectfully submitted,

/DAVID L. VANIK/
David L. Vanik, Ph.D.
Registration No.: 64,547

Susan E. Shaw McBee
Registration No. 39,294